Attribution

This report is based on the original work of an expert panel led by Gianfranco Pezzino, M.D., M.P.H. and Steven Q. Simpson, M.D.

Members of the panel are listed in Appendix A. The project was conducted in 2009-2010 by the Kansas Health Institute under contract with the Kansas Department of Health and Environment.
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BACKGROUND

These guidelines describe principles and practices that health care providers and acute care hospitals in Kansas should adopt if resources are scarce during a public health emergency. The guidelines are the product of analysis conducted on behalf of the Kansas Department of Health and Environment (KDHE) by the Kansas Health Institute (KHI) and a panel of experts. In September 2009, KHI produced a report for KDHE that outlined possible general processes and ethical principles to apply when health care resources are scarce. The report also recommended that KDHE develop and distribute as soon as possible to providers guidelines that address potential lack of resource situations that may occur as a result of the current influenza pandemic.

In response to the report, KDHE asked KHI to convene a group of medical experts to review and amend, as deemed necessary, four technical documents developed by other states that the report identified as good references on the subject. In May, 2013 a re-analysis was undertaken by the KDHE’s Clinical Resource Network (CRN) and updates were made for release in September, 2013. The guidelines presented in this document represent the result of the expert panel review and of comments received from health care professionals. The names and affiliation of the members of the review panel are listed in Appendix A of this document.

Since this document’s recommended guidelines are adapted from previous publications, the background information and detailed rationale for the guidelines have been considerably shortened since the review panel’s goal was to produce a concise list of recommendations that clinicians in Kansas could rapidly review and implement. Those interested in more background information and further justification of the guidelines can review the original source documents, which are listed below:


“Summary of Suggestions from the Task Force for Mass Critical Care Summit”, January 26–27, 2007, [http://www.chestjournal.org/content/133/5_suppl/1S.full.pdf+html].


1 The KHI report can be found at: www.khi.org
GENERAL PRINCIPLES

• These modified protocols of care should not be considered a substitute for good planning of regional sharing of resources and surge capacity. The activation of the modified protocols should take place only after a declaration of emergency and only after other specified means of procuring additional resources and expanding surge capacity have been exhausted.

• These protocols address primarily hospital triage and should be integrated into broader emergency response plans. For example, the adoption of these protocols could require that some patients be moved after triage to reference hospitals to receive life-saving treatment or out of acute care hospitals if they do not qualify for life-saving treatment. This and similar issues should be addressed in local and state emergency response plans.

• Hospitals should work within the framework of regional networks, i.e. the Kansas Preparedness Healthcare Coalitions that are already in place. Resource deficiency may be a local or regional problem and could be mitigated by carefully drafted mutual aid and sharing protocols. Regional networks could also play a vital role in assuring that the modified protocols can be implemented throughout the state, with small and large hospitals working together to assure a uniform process of triage and allocation of resources.

• Before these modified protocols are implemented, all key stakeholders should be aware of the specifics to ensure that there is sufficient clarity and consensus to implement them.

• Small hospitals may have difficulty adopting some of the modified protocols proposed in this document. The review panel discussed this issue and concluded that, while modified protocols that provide for the same solution for all may not be always easy to implement, they have the advantage of promoting a fairer and more uniform distribution of resources throughout the state. When applicable, specific differences in implementation between small and large hospitals and communities are addressed and discussed in the protocols. Additional adjustments may be necessary based on new experiences and evidence. Issues concerning small hospitals are discussed further in a special section of this document.

• Because the field of modified protocols of care is so new, and interventions have not been widely tested, the panel strongly recommends that all the protocols be labeled as “Interim Recommendations.” This will facilitate changing and updating the documents as new information becomes available.

• The panel recommends that KDHE issue the protocols as voluntary, not mandatory, guidelines. The panel expects that the declaration of emergency that would trigger the implementation of these protocols would also offer liability protection under the provisions of K.S.A. 48-915 (b) to health care providers and hospitals that implement them in good faith. The panel trusts that such protection will help remove any reservations that institutions and clinicians might have about implementing these protocols, allowing a broader, and therefore more effective, implementation of the protocols.
SPECIAL ISSUES CONCERNING SMALL HOSPITALS

Small hospitals may have difficulty adopting some of the modified protocols proposed in this document. The review panel discussed this issue and tried to leave as much flexibility as possible in the protocols to account for local circumstances, while assuring a standardized approach to the use of scarce resources throughout the state. Modified protocols that provide for the same solution for all situations may not be easy to implement, but they promote a fairer and more uniform distribution of resources throughout the state. During the comment period, questions were raised about the feasibility of implementing the modified protocols in small hospitals; but no evidence surfaced suggesting that implementation in small hospitals would not be possible through careful planning and via the regional networks.

When small hospitals do not have the resources to triage or treat patients locally using the proposed modified protocols, we recommend that they work in close partnership with their referral institutions. It could be possible, for example, to appoint a triage officer in a large hospital who could conduct triage for patients admitted in a small hospital. The triage could be conducted remotely using teleconferences or, if necessary, telemedicine resources. It is important that triage decisions for critically ill patients occur at the local level, even if the decisions are made by a triage officer in a different institution. As one of the providers told us, “there is no sense in transferring patients who will be very low priority patients when they arrive at the referral center.” Some of these mechanisms of assisted remote triage may already be in place and used occasionally during localized emergencies.

Large hospitals should be ready to assist small hospitals with their triage needs, and to treat their patients and patients transferred from small hospitals using the same set of clinical priority criteria. In the absence of this uniform approach, it is likely that patients in rural areas and those closer to referral hospitals would be treated unequally, creating a situation of geographical disparity that would be in contrast with the principles of distributive justice endorsed in this document. Such a situation could also create uncontrolled movement of patients towards large hospitals, in the hope that they could be treated there, which would increase congestion in those institutions. To obviate such a one way flow of patients, it may be necessary for larger, referral facilities to send less critically ill patients, who are not requiring the specialized capabilities of the referral center, to the smaller hospitals for ongoing care and completion of hospitalization.

The adoption of clinical triage criteria specific to small hospitals also was examined. In particular, the use of a modified SOFA score that uses saturation of peripheral oxygen (SpO2) instead of partial pressure of oxygen in arterial blood (PaO2) was considered, since some hospitals do not
perform the Arterial Blood Gas analysis test (ABG) necessary to measure $\text{PaO}_2$. In the absence of convincing published evidence in support of the modified SOFA score the review panel decided to endorse the use of the unmodified SOFA criteria throughout the state. The panel recommends that hospitals review the requirements for the SOFA assessment and make provisions to assure that they have the capacity to perform the necessary laboratory tests.
APPENDIX A: EXPERT PANEL MEMBERS

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Gianfranco Pezzino, MD, MPH, Senior Fellow, Kansas Health Institute, Topeka, KS

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EXPERT PANEL MEMBERS

Original Version of Modified Health Care Protocols

Originally Published by the Kansas Health Institute, November, 2009; Revised, August 2010

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APPENDIX B: INTERIM GUIDELINES FOR TERTIARY TRIAGE PROTOCOL FOR ALLOCATION OF SCARCE RESOURCES IN ACUTE CARE HOSPITALS IN KANSAS

I. GOAL

1. This protocol should be used in hospitals throughout Kansas to ensure that patients have equitable access to life-saving resources when the demand for these resources is greater than the supply, and when use of resources must be optimized.

2. The application of these guidelines in small hospitals may not be feasible due to the lack of specialized staff. In these cases, hospitals may modify the implementation of these guidelines to fit their situation while preserving the overarching goal of assuring an objective, clinical set of criteria for the allocation of scarce resources. Small hospitals should also partner with larger referral centers and delegate some functions described in this document to those centers. Communication between small and large hospitals can take place using the best and most appropriate means, such as telephone, radio, telemedicine, or face-to-face consultation.

3. While the protocol refers primarily to pandemic influenza, it is applicable to other public health emergencies that may cause a prolonged shortage of life-saving resources, such as chemical disasters, tornado or other weather-induced disasters, or acts of terrorism.

II. INITIATION OF THE TRIAGE PROTOCOL

1. Generally, the hospital medical director, in consultation with the hospital administrator, will apply the protocol throughout an affected hospital at his or her discretion. The medical director will take into consideration local or regional declarations of emergency (e.g., state-wide declaration of emergency by the governor).

2. Hospital medical directors must assure that the protocol is applied consistently and fairly whenever and wherever it is initiated.

3. Application of the pandemic triage protocol will take place only when augmentation efforts have been exhausted and demand for the life-saving resource exceeds supply. Triggers include (but are not limited to):

   a. Local or state declaration of emergency.

   b. Initiation of national disaster medical system and national mutual aid and resource management.

   c. Surge capacity fully employed within health care facility

   d. Attempts at conservation, reutilization, adaptation, and substitution are performed maximally

   e. Identification of critically limited resources (ventilators, antibiotics)

2 Last revised: August 9, 2010
f. Request for resources and infrastructure made to local and state health officials

g. Current attempt at regional, state, and federal level for resource or infrastructure allocation

4. The hospital medical director should rescind the application of the pandemic triage protocol when the supply of the life-saving resource is sufficient to meet the demand. This may occur either before or after a declared state of emergency has been rescinded.

III. RESPONSIBILITY STRUCTURE FOR TRIAGE DECISION MAKING

1. Scarce Resource Allocation Team:

   a. The scarce resource allocation team should be a functional team under existing Incident Command System (ICS)/Hospital Incident Command System (HICS)/Emergency Operations — it should not be a separate structure.

   b. The size and composition of the allocation team will vary depending on local circumstances, the nature of the emergency, and the size of the institution. Members may include (but not be limited to) critical care physicians, critical care nurses, respiratory therapists, pharmacists, human resource managers, hospital administrators and legal counsel.

   c. The scarce resource allocation team will:

      i. Acquire the information necessary to facilitate and oversee informed and ethical triage and scarce resource allocation decisions. Information could include resources (bed census, staffing, projected needs for care, existing medical resources, resource gaps, and projected availability of life-saving and hospice and palliative care resources) and guidelines for the management of the emergency (e.g., up-to-date treatment options and prognostic factors).

      ii. As part of Incident Command System (ICS)/Hospital Incident Command System (HICS)/Emergency Operations, make judgments in collaboration with health care organization leaders and staff to implement appropriate alternative standard protocols of care that address the special demands that an emergency imposes on the health care organization or demands that could imminently be expected.

      iii. Meet often, at least daily, during an emergency.

      iv. Advise and assist, as required, and make definitive decisions, if necessary, to resolve uncertainties and disputes that affect the health care organization’s capacity to carry out its mission during a public health emergency.

      v. Be involved in the real-time appeals process regarding triage decisions described in this document (excluding decisions made by members of the triage team which should not be subject to appeal).

      vi. Prepare information briefs to the chief executive officer, chief of staff or designee(s) about the emergency’s status and the health care organization’s response so that the information may be communicated to appropriate staff and stakeholders.
2. Triage Officer:

a. The triage officer must be a qualified member of the medical staff who is, ideally, experienced and trained in intensive care and triage protocols.

b. The triage officer will assess all patients; assign a level of priority for each, and direct attention to the highest-priority patients.

c. The triage officer, with the assistance of the triage team (when available), will:

   i. Review all patients for inclusion and exclusion criteria, and facilitate discharge from critical care for patients no longer requiring it.

   ii. At least every 24 hours, evaluate all patients receiving critical care.

   iii. Evaluate all patients that have been recommended to receive critical care.

d. The triage officer is not expected to examine patients, except under circumstances in which examination may be crucial in reaching a triage decision.

e. The triage officer should not be involved in day to day care of the patients subjected to triage. Small hospitals unable to maintain this separation of roles should use a triage officer based in another institution. Such individuals may be identified by reference to the Regional Healthcare Coalition documents. Each hospital should pre-identify potential individuals for off-site triage for use in the event of disaster circumstances.

f. The triage officer will make triage decisions based on the allocation protocol, assigning patients to triage categories based on a SOFA score or exclusion criteria (Tables 2 and 3), and on available resources.

3. Triage Team:

a. In hospitals with sufficient staff resources, a triage team will be set up as a subcommittee of the scarce resource allocation team.

b. The role of the triage team is to provide information to the triage officer and help facilitate and support his or her decision-making process.

c. Members of the triage team may include (but not be limited to) an experienced critical care nurse, respiratory therapist, or clinical pharmacist. A representative from hospital administration may also be a part of the team to help organize resources and serve as a liaison to hospital leadership.

d. In larger facilities, it may be necessary to have more than one triage officer and team, with each officer/team assigned to a designated ICU or hospital area and to specific operational periods or shifts. In such circumstances, triage personnel should designate time for mutual review and transition of ongoing triage issues. It is recommended that the triage officer and team members function in shifts lasting no longer than 12 to 16 hours, if feasible.
e. The triage officer and triage team will:

i. Meet often (at least daily) to assess all patients who have clinical indications to receive scarce life-saving resources (e.g., critical care patients who require ventilators or hemodynamic support) and evaluate exclusion and inclusion criteria to determine the appropriateness of the initiation and continuation of scarce life-saving treatment.

ii. Develop and maintain a record of triage decisions including the data upon which the decisions were based.

f. Decisions from the triage team/triage officer cannot be appealed.

4. Review Committee:

a. In hospitals with sufficient staff resources, a review committee will be created to review the decisions of the triage team.

b. The review committee (ideally a small group of no more than three individuals) may be composed of experienced professionals who typically no longer provide direct care, such as the chief nursing officer, chief medical officer, chief respiratory therapy supervisor, infection control director, or legal counsel.

c. The review committee will bring to the attention of the triage officer any concerns about the application of the triage algorithm so that the triage officer may reflect on these concerns when approaching future decisions.

d. The review committee does not have the authority to change a decision made by the triage officer, except when there is clear evidence that the triage protocol was not applied as planned.

5. Treating Clinicians:

a. Should not have the responsibility of deciding whether to institute or remove a patient from life-saving resources. This decision is up to the triage team/triage officer. These functions should be kept separated to reduce the emotional impact of these decisions on health care providers.

b. Will implement a treatment plan consistent with the triage team’s decision regarding patient triage category.

c. Will conduct a DNR discussion with patients who do not qualify under the triage protocol for scarce life-saving resources.

d. Will offer palliative and other appropriate care.

6. Emergency Physicians:

a. Because many patients will seek care at the emergency department during pandemic influenza, emergency department personnel should be prepared to apply the “initial assessment tool” (See Table 3) for patients who have clinical indications for critical care.
b. Emergency physicians will:
   i. Apply initial resuscitation, if applicable, with simple measures such as fluids oxygen by nasal cannula, mask, and control of bleeding, etc. (unless other exclusion criteria are present).
   ii. Report initial assessment to the triage team.

IV. ALLOCATION CRITERIA

1. The overarching criterion is the degree of medical success or survivability determined by the application of established, objective clinical criteria, including SOFA scores. The guiding question of this assessment is whether the patient is likely to survive with the use of the scarce resource.

2. Once a determination has been made that a patient qualifies for the resource under the SOFA score, and a patient’s priority category has been determined, within-category priority will be established on a first-come, first-served basis or on a random selection/lottery basis, depending on feasibility of implementation.
   a. This second step will be implemented only if resources are still insufficient to meet the needs of all who qualify for the resource, after applying the clinical allocation criteria.

3. Clinical Assessment
   a. Clinicians will thoroughly assess all patients who present for care.
   b. Patients with clinical indications for scarce life-saving resources (e.g., critical care patients who require ventilators or hemodynamic support) will be subject to the triage protocol described in this document, unless they elect not to be candidates for critical care.

4. Exclusion Criteria
   a. Patients with clinical indications for scarce life-saving resources will be assessed for exclusion criteria to determine the appropriateness of the initiation or continuation of scarce life-saving treatment.
   b. Exclusion criteria are intended to identify and exclude patients with a short life expectancy irrespective of the current acute illness. If an exclusion criterion is present (Table 1), the patient is no longer a candidate for scarce life-saving resources, including scarce resources that may be needed for cardiopulmonary resuscitation.
   c. Clinicians should offer palliative and other supportive care to the patient and follow clinical standards for withdrawal of scarce life-saving resources.

V. RE-ASSESSMENT

1. Continued use of the scarce life-saving resources will be reviewed on an established

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3 The triage of patients with a Do Not Resuscitate (DNR) order or other advance directives should take into account the patient’s wishes and the likelihood of recovery after life-sustaining measures are applied.
schedule by the triage team (at least once every 24 hours). Patients that continue to meet
criteria for inclusion will receive the resources until they either meet an exclusion
criterion, or they are re-assessed according to the triage team schedule.

a. Patients assigned to the same category will be allocated resources on a first-come,
first-served basis or on a random selection/lottery basis, depending on the feasibility
of implementation.

b. Those that no longer meet the criteria after re-assessment will no longer be eligible
for access to the scarce life-saving resources and should be informed of the need for
withdrawal of these treatments.

VI. SPECIAL CONSIDERATIONS FOR VENTILATORS

1. Allocation of ventilators during a public health emergency will be subject to the same
procedures described in this document for other scarce resources. Since ventilators are
often an important life-saving resource, this section reviews some special issues related to
ventilator allocation. For more details please refer to the following document, from which
many of these guidelines have been abstracted:

“NYS Workgroup on Ventilator Allocation in an Influenza Pandemic. Allocation of
Ventilators in an Influenza Pandemic”, March 15, 2007,
ventilator_guidance.pdf]

2. Uniform policies are crucial; variations among facilities will lead to inequities. Equitable
rationing systems, particularly ones that contemplate limiting access to lifesaving
treatment, must assure that the same resources are available and in use at similarly
situated facilities, i.e., all facilities in one city gripped by the pandemic.

3. The establishment of regional stockpiles should be strongly considered, following the
example in New York and other states. Leaders of facilities within a region should
be encouraged to work out voluntary plans for loans of equipment and staff in a crisis.

4. As a public health emergency spreads, hospitals should limit the non-critical use of
ventilators. Elective procedures that may require the use of ventilators should be canceled
or postponed during the period of emergency. For an emergency that stretches from days
to weeks, such as a pandemic, facilities will need a review system for procedures that
decrease morbidity or mortality, but are not of an emergency nature.

5. The ideal interval for re-assessing patients in need of critical care and ventilators has not
been well defined. Critical care experts point out that many patients will not show signs
of improvement for several days after they start receiving intensive care resources such as
ventilators; therefore a re-assessment schedule should allow for sufficient time to pass
from when a patient first receives the resources, so that clinical improvement can become
evident. Other experts point out that the greatest impact on survival is often made by
aggressive action in the first hours of presentation, and a reassessment schedule that is
conducted using long intervals may not identify early enough patients who fail to
improve (and whose critical care resources should therefore be re-allocated). These are
factors that should be kept in mind when determining a re-assessment schedule. The
decision should be based on the clinical characteristics of the emergency and on how
acute the need for the re-allocation of resources is. The expert panel believes that
hospitals should reassess this allocation every 24 hours.
6. Distinctions should be maintained between acute and chronic care facilities once triage begins, permitting chronic care facilities to maintain their specific mission. Patients using ventilators in chronic care facilities would not be subjected to acute care triage guidelines. If, however, such patients required transfer to an acute care facility, they would be assessed by the same criteria as all other patients, and might fail to meet criteria for continued ventilator use. Chronically ill patients will be vulnerable to the pandemic; chronic care facilities will have to provide more intensive care on site as part of the general process of expanding care beyond standard locations. Barriers to transfer are appropriate and likely during a phase in which acute care hospitals are overwhelmed.

7. Children in need of ventilators present unique challenges.

a. In general, triage using SOFA scores should not be used for children (especially young ones), because the SOFA system has not been adequately tested in children.

b. The use of the modified system described in Appendix C of this document (*Interim Guidelines for the Use of Pediatric Ventilators During a Public Health Emergency in Kansas*) is recommended in alternative to the SOFA triage system for children.

c. Special expertise, likely to be in short supply, is needed to care for children who may also be especially vulnerable to morbidity and mortality in a pandemic. The establishment of centers of excellence for pediatric patients, particularly during a pandemic, should be considered. Although a pandemic emergency is likely to affect most or all of the state, the required expertise will not be widely distributed and an attempt to concentrate severely ill children needing of intensive care in specialized centers may make sense, if feasible. Transportation of pediatric patients to the referral centers may be problematic in the middle of a statewide emergency, when the emergency medical system could be under considerable pressure.

d. Planning assumptions must adequately reflect the needs of infants and children. Many modern ventilators accommodate patients weighing as little as 10 kilograms, but will not support infants.
**Table 1. Exclusion Criteria**

<p>| Severe, advanced chronic disease with a short life expectancy (6 months or less) |
| Severe burns on patient with any two of the following: |
| Age &gt; 60 yr |
| 40% of total body surface area affected |
| Inhalational injury |
| Cardiac arrest: |
| Un-witnessed cardiac arrest |
| Witnessed cardiac arrest, not responsive to electrical therapy (defibrillation or pacing) |
| Recurrent cardiac arrest Trauma-related arrest |
| Advanced untreatable neuromuscular disease |
| Metastatic malignant disease with poor prognosis |
| End-stage organ failure (except when caused by readily reversible volume overload or hypoventilation due to an exogenous agent, such as narcotic, benzodiazepine, or other procedural sedative): |
| Cardiac: NY Heart Association class III or IV |
| Pulmonary: severe chronic lung disease with FEV1** &lt; 25% |
| Hepatic: MELD*** score &gt; 20 |
| Renal: dialysis dependent |
| Neurologic: severe, irreversible neurologic event/condition with high expected mortality |</p>
<table>
<thead>
<tr>
<th>Variable</th>
<th>SOFA Score</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0</td>
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<tr>
<td>PaO2/FiO2 mmHg</td>
<td>&gt; 400</td>
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<tr>
<td>Platelets, x 103/μL or x 106/L</td>
<td>&gt; 150</td>
</tr>
<tr>
<td>Bilirubin, mg/dL (μmol/L)</td>
<td>&lt; 1.2 (≤ 20)</td>
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<td>Hypotension</td>
<td>None</td>
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<tr>
<td>Glasgow Coma Score</td>
<td>15</td>
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<tr>
<td>Creatinine, mg/dL (μmol/L)</td>
<td>&lt; 1.2 (≤ 106)</td>
</tr>
</tbody>
</table>

Note: Clinicians will determine the total SOFA score for each patient by summing the scores for each variable. Dopamine [Dop], epinephrine [Epi], norepinephrine [Norepi] doses in ug/kg/min. SI units are noted in parentheses ( ).

*Adapted from: Ferreira et al., 2001. Explanation of variables: PaO₂/FiO₂ indicates the level of oxygen in the patient’s blood. Platelets are a critical component of blood clotting. Bilirubin is measured by a blood test and indicates liver function. Hypotension indicates low blood pressure; scores of 2, 3, and 4 indicate that blood pressure must be maintained by the use of powerful medications that require ICU monitoring, including dopamine, epinephrine, and norepinephrine. The Glasgow coma score is a standardized measure that indicates neurologic function; low score indicates poorer function. Creatinine is measured by a blood test and indicates kidney function.

<table>
<thead>
<tr>
<th>Initial Criteria</th>
<th>Priority</th>
<th>Action</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exclusion Criteria</td>
<td>None</td>
<td>Do not use life-saving resources</td>
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<tr>
<td>OR</td>
<td></td>
<td>Use other resources including palliative measures</td>
</tr>
<tr>
<td>SOFA &gt; 11</td>
<td></td>
<td></td>
</tr>
<tr>
<td>SOFA &lt; 7</td>
<td>Highest</td>
<td>Use life-saving resources, as available</td>
</tr>
<tr>
<td>Single Organ Failure</td>
<td>Intermediate</td>
<td>Use life-saving resources, as available</td>
</tr>
<tr>
<td>SOFA 8–11</td>
<td>Intermediate</td>
<td>Use life-saving resources, as available</td>
</tr>
<tr>
<td>No requirement for life-saving resources</td>
<td>None</td>
<td>Use other medical management</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Re-assess as needed</td>
</tr>
<tr>
<td>48 Hour Criteria</td>
<td>Priority</td>
<td>Action</td>
</tr>
<tr>
<td>------------------------------------------------------</td>
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<td>------------------------------------------------------------------------</td>
</tr>
<tr>
<td>Exclusion Criteria</td>
<td>None</td>
<td>Discontinue life-saving resources</td>
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<tr>
<td><strong>OR</strong></td>
<td></td>
<td>Use other resources including palliative measures</td>
</tr>
<tr>
<td>SOFA &gt; 11</td>
<td></td>
<td></td>
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<tr>
<td><strong>OR</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>SOFA 8 – 11 and increasing since last assessment</td>
<td></td>
<td></td>
</tr>
<tr>
<td>SOFA &lt; 11 and decreasing since last assessment</td>
<td>Highest</td>
<td>Continue life-saving resources, as available</td>
</tr>
<tr>
<td>SOFA &lt; 11 and unchanged since last assessment</td>
<td>Intermediate</td>
<td>Continue life-saving resources, as available</td>
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<tr>
<td><strong>OR</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>SOFA &lt; 8 and increasing since last assessment</td>
<td></td>
<td></td>
</tr>
<tr>
<td>No longer requiring life-saving resources</td>
<td>None</td>
<td>Discontinue life-saving resources. Re-assess as needed</td>
</tr>
</tbody>
</table>

* Re-assessment should be conducted on a predetermined scheduled, at least every 24 hours.
# APPENDIX C: INTERIM GUIDELINES FOR THE USE OF PEDIATRIC VENTILATORS DURING A PUBLIC HEALTH EMERGENCY IN KANSAS

## PELOD Scoring System

<table>
<thead>
<tr>
<th>Organ System</th>
<th>Variable</th>
<th>0</th>
<th>1</th>
<th>10</th>
<th>20</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Neurologic</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Glasgow coma score</td>
<td>12-15</td>
<td>7-11</td>
<td>4-6</td>
<td>3</td>
</tr>
<tr>
<td></td>
<td>AND</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>OR</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Papillary reaction</td>
<td>Both reactive</td>
<td>Both fixed</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Cardiovascular</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Heart rate</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>&lt;12 y</td>
<td>≤195 bpm</td>
<td>&gt;195 bpm</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>&gt;12 y</td>
<td>≤150 bpm</td>
<td>&gt;150 bpm</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>AND</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>OR</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Systolic blood pressure</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>&lt;1 mo</td>
<td>&gt;65 mm Hg</td>
<td>35-65 mm Hg</td>
<td>&lt;35 mm Hg</td>
<td></td>
</tr>
<tr>
<td></td>
<td>≥1 mo &amp; &lt; 1 yr</td>
<td>&gt;75 mm Hg</td>
<td>35-75 mm Hg</td>
<td>&lt;35 mm Hg</td>
<td></td>
</tr>
<tr>
<td></td>
<td>≥1 yr &amp; &lt; 12 y</td>
<td>&gt;85 mm Hg</td>
<td>45-85 mm Hg</td>
<td>&lt;45 mm Hg</td>
<td></td>
</tr>
<tr>
<td></td>
<td>≥12 y</td>
<td>&gt;95 mm Hg</td>
<td>55-95 mm Hg</td>
<td>&lt;55 mm Hg</td>
<td></td>
</tr>
<tr>
<td><strong>Renal</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Creatinine</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>&lt;7d</td>
<td>&lt;1.59 mg/dL</td>
<td>≥1.59 mg/dL</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>≥7d &amp; &lt; 1 y</td>
<td>&lt;0.62 mg/dL</td>
<td>≥0.62 mg/dL</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>≥ 1 y &amp; &lt; 12 y</td>
<td>&lt;1.13 mg/dL</td>
<td>≥1.13 mg/dL</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>≥12 y</td>
<td>&lt;1.59 mg/dL</td>
<td>≥1.59 mg/dL</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Pulmonary</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Pa O₂/F₁₀₂ ratio</td>
<td>&gt;70 mm Hg</td>
<td>≤70 mm Hg</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>AND</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>OR</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Pa CO₂</td>
<td>≤90 mm Hg</td>
<td>&gt;90 mm Hg</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>AND</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Mechanical vent</td>
<td>No</td>
<td>Yes</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Hematologic</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>WBC</td>
<td>≥4.5 K</td>
<td>1.5-4.4 K</td>
<td>&lt;1.5</td>
<td>10</td>
</tr>
<tr>
<td></td>
<td>AND</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>OR</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Platelets</td>
<td>≥35 K</td>
<td>&lt;35</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Hepatic</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>AST</td>
<td>&lt;950 IU/L</td>
<td>≥950 IU/L</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>AND</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Prothrombin time</td>
<td>&gt;60%</td>
<td>≤60%</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

1 Abbreviations: PELOD, Pediatric Logistic Organ Dysfunction; bpm, blood pressure monitor; Pa O₂/F₁₀₂, partial pressure of oxygen, arterial/fraction of inspired oxygen; Pa CO₂, partial pressure of carbon dioxide, arterial; WBC, white blood cells; AST, aspartate aminotransferase.

Development of a Pediatric Multiple Organ Dysfunction Score: Use of Two Strategies

Stéphane Leteurtre, Alain Martinot, Alain Duhamel, France Gauvin, Bruno Grandbastien, Thi Vu Nam, François Proulx
Critical Care Triage Tool - Pediatric Patients (<18y) (Top), and Exclusion Criteria (Bottom).

<table>
<thead>
<tr>
<th>Color Code</th>
<th>Initial Assessment</th>
<th>48-h Assessment</th>
<th>120-h Assessment</th>
</tr>
</thead>
</table>
| Blue       | Exclusion criteria  
OR
PELOD ≥ 33 | Medical management +/- palliate & discharge from critical care | Exclusion criteria  
OR
PELOD ≥ 33 or PELOD 21-33 & no change | Palliate & discharge from critical care |

| Red        | PELOD ≤ 21  
OR
Single organ failure | Highest | PELOD < 33 and decreasing | Highest | PELOD < 33 and decreasing progressively | Highest |

| Yellow     | PELOD 21-33 | Intermediate | PELOD < 21 no change | Intermediate | PELOD < 21 minimal decrease (<3-point decrease in past 72 h) | Intermediate |

| Green      | No significant organ failure | Defer or discharge, reassess as needed | No longer ventilator dependent | Discharge from critical care | No longer ventilator dependent | Discharge from critical care |

**Exclusion Criteria**

Patient is excluded from admission or transfer to critical care if any of the following is present:

- A Sever trauma
- B Severe burns of patient with any two of the following:
  - Age > 60 y
  - >40% of total body surface area affected
  - Inhalation injury
- C Cardiac arrest
  - Unwitnessed cardiac arrest
  - Witnessed cardiac arrest, not responsive to electrical therapy (defibrillation or pacing)
  - Recurrent cardiac arrest
- D Metastatic malignant disease with poor prognosis
- E Advanced and irreversible immunocompromise
- F Severe and irreversible neurologic event or condition with highly expected mortality
- G End-stage organ failure meeting the following criteria:
  - **Heart**
    - NYHA class III or IV heart failure
  - **Lungs**
    - Severe chronic lung disease with FEV\(_1\) < 25% predicted, baseline PaO\(_2\) < 55 mm Hg, or secondary pulmonary hypertension
    - Previously diagnosed primary pulmonary hypertension with NYHA class III or IV heart failure, or mean pulmonary arterial pressure > 50 mm Hg
  - **Liver**
    - Child-Pugh score > 7 or MELD score of > 20

NYHA New York Heart Association; FEV\(_1\) forced expiratory volume in the first second of expiration; MELD, model for end-stage liver disease
Core strategies that can be employed (generally in order of preference) during or in anticipation of a scarce resource situation are:

**Prepare**—pre-event actions taken to minimize resource scarcity.

**Substitute**—use an essentially equivalent device, drug, or personnel for one that would usually be available (e.g., morphine for fentanyl).

**Adapt**—use a device, drug, or personnel that are not equivalent but that will provide sufficient care (e.g., anesthesia machine for mechanical ventilation).

**Conserve**—use less of a resource by lowering dosage or changing utilization practices (e.g., minimizing use of oxygen driven nebulizers to conserve oxygen).

**Re-use**—re-use (after appropriate disinfection / sterilization) items that would normally be single-use items.

**Re-allocate**—take a resource from one patient and giving it to a patient with a better prognosis or greater need.

Examples of the application of these strategies are presented below. Some examples refer to situations that may take place outside of a public health emergency and may already be addressed by medical staff.

### Oxygen

**Conserve strategy**—Use minimum liter flow to keep O2 saturation > target (85–95% depending on situation). Use O2 conserving cannulas (Oxymizer™). No oxygen driven nebs. Eliminate or reduce equipment with high O2 consumption.

**Re-Use strategy**—Appropriately disinfect and re-use cannulas, masks, and tubing.

**Re-Allocate strategy**—May have to base therapy on triage decision tool similar to ventilator allocation.

### Medication Administration

**Substitute strategy**—Use alternative inexpensive medications (morphine, lorazepam, doxycycline) that are easily stockpiled prior to the event.

**Adapt strategy**—Use morphine and benzodiazepines for sedation drips, when possible. Run drips via gravity rather than IV pump, if needed. Administer more medications via a subcutaneous or
intramuscular route rather than intravenously.

**Conserve strategy**—Give adjunctive non-steroidal and other analgesics/medications including orally when possible.

**Re-Allocate strategy**—Re-allocation should be considered as the last resort. Re-allocation will increase demands for palliative care and adequate pain control/sedation—focus should be on stockpiling inexpensive options in advance of event.

**Hemodynamic Support and IV Fluids**

**Substitute strategy**—Use alternative vasopressor agents such as epinephrine (inexpensive).

**Adapt strategy**—May have higher threshold to initiate vasopressors, may use gravity drips (e.g., 1mg epinephrine in 100cc NS) instead of infusion pumps. Consider nasogastric fluid replacement rather than IV.

**Conserve strategy**—Minimize invasive monitoring.

**Re-Use strategy**—Consider reusing central venous catheters, other tubes and catheters with appropriate sterilization/disinfection.

**Mechanical Ventilation**

**Adapt strategy**—Use of anesthesia machines, BiPAP, short-term manual ventilation and other strategies.

**Conserve strategy**—Adjusted threshold for intubation, decrease elective surgeries to free up ventilators/anesthesia machines.

**Re-Use strategy**—Re-use of ventilator circuits after appropriate sterilization / disinfection.

**Re-Allocate strategy**—Re-allocation should be considered as the last resort. Ventilators should be allocated to patients who can most benefit, and allocation should follow a pre-planned process and use decision support tools and expert clinical judgment.

**Nutrition**

**Adapt strategy**—Have family or ancillary staff provide meals. Provide simpler meals and offer fewer choices to those that can take oral intake. Use tube feedings instead of total parenteral nutrition when possible. Delay feedings longer than usual.

**Conserve strategy**—See above.

**Re-Use strategy**—May need to re-use nasogastric and other feeding equipment with appropriate disinfection.

**Staffing**

**Substitute strategy**—Outside, equally-qualified staff brought in to institution via compact agreements or other mechanism (DMAT, Medical Reserve Corps, other local/regional/state/federal sources). Use family or non-professional staff to provide basic patient cares (non-clinical).
Adapt strategy—Less qualified staff from sources as above or volunteers provide basic patient care with critical care nursing and physician staff monitoring larger numbers of patients. Implement just-in-time training and orientation to job duties following pre-planned training programs. Change shift duration. Use family or non-professional staff to provide some clinical care with training/in-service.

Conserve strategy—Reduce administrative demands (teaching and administration, documentation, etc.).